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~~3~~ ~~17~~. (New) The method of claim ~~16~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is administered prophylactically or therapeutically to prevent or treat peritoneal adhesions.

cont
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²
~~4~~ ~~18~~. (New) The method of claim ~~16~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is administered by intraperitoneal injection of a composition comprising the bioactive glass particles, a suitable carrier for intraperitoneal injection, and one or more therapeutic agents.

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~~5~~ ~~19~~. (New) The method of claim ~~18~~ wherein the one or more therapeutic agents are selected from the group consisting of healing promotion agents, growth factors, anti-inflammatory agents, and anesthetics.

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~~6~~ ~~20~~. (New) The method of claim ~~16~~ wherein the bioactive glass particles have a size less than about 2 microns.

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~~7~~ ~~21~~. (New) The method of claim ~~12~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is administered by inhalation.

¹
~~8~~ ~~22~~. (New) The method of claim ~~12~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is administered by subcutaneous injection.

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~~9~~ ~~23~~. (New) The method of claim ~~22~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is mixed with a biocompatible hydrogel.

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~~10~~ ~~24~~. (New) The method of claim ~~22~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is administered at a site at which surgery is to be performed.

¹⁰
~~11~~ ~~25~~. (New) The method of claim ~~24~~ wherein the locally effective TNF- α lowering amount of bioactive glass particles is mixed with an anesthetic.
